

Division: Pharmacy Policy	Subject: Prior Authorization Criteria
Original Development Date: Original Effective Date:	February 9, 2016
Revision Date:	February 24, 2016; January 21, 2022; June 16, 2022, January 9, 2023, May 11, 2023, January 24, 2024

Testosterone (non-injectable formulations)

TOPICAL:

- Clinical PA required (preferred): Androderm[®] patch, Testim[®] gel tube, Testosterone gel packet (Vogelxo), Testosterone gel pump (AndroGel[®])
- Clinical PA required (non-Preferred): AndroGel[®] pump, Fortesta[®] gel pump, Natesto[®] nasal gel pump, Testosterone gel packet/tubes, Testosterone Pump (Axiron[®]), Testosterone topical solution, Vogelxo[®] gel packet/pump/tube

IMPLANT:

o Preferred: N/A

Non-Preferred: Testopel[®]

ORAL:

Preferred: N/A

o Non-Preferred: Kyzatrex[®], Jatenzo[®], Tlando[®]

LENGTH OF AUTHORIZATION: One year

INITIAL REVIEW CRITERIA:

- Patient is \geq 18 years old; AND
- Patient is male; AND
- Patient has a diagnosis of primary or secondary hypogonadism;* AND
- Patient does not have a history of prostate carcinoma or male breast carcinoma; AND
- Prescriber has submitted the results of two separate serum testosterone levels, each drawn in themorning, which indicate a low serum testosterone (normal range: 300 to 1,000 ng/dL) within thelast six months.
- * Causes of hypogonadism are classified as primary which are due to failure of the testes, or secondary, which are due to failure of the hypothalamus or pituitary gland. Either type of hypogonadism, may be caused by an inherited (congenital) or acquired factor.
- * Examples of primary male hypogonadism include but are not limited to cryptorchidism, bilateral torsion, orchitis, vanishing testis syndrome, orchidectomy, chemotherapy, radiation therapy, toxic damagefrom alcohol or heavy metals, testicular infections (such as mumps) and chromosomal abnormalities such as Klinefelter's Syndrome *Examples of secondary male hypogonadism include but are not limited idiopathic gonadotropin orluteinizing hormone releasing hormone (LHRH) deficiency and pituitary hypothalamic injury fromtumors, trauma, or radiation.

PATIENTS WHO MEET CRITERIA SHOULD BE APPROVED FOR THE PREFERRED AGENTS



^{**}Safety and efficacy in men "age-related hypogonadism" has not been established.



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CONTINUATION OF THERAPY CRITERIA:

- Patient has been compliant with treatment based on refill history
- Prescriber submits labs within the last twelve months indicating patient has a normal serum testosterone level on therapy (normal range: 300-1,000 ng/dL)

DOSING & ADMINISTRATION:

Refer to product labeling at https://www.accessdata.fda.gov/scripts/cder/daf/

